

TITLE: Purification and properties of L-  
 asparaginase produced by *Aspergillus niger*, S-48  
 TAT, the causal fungus of  
 biodeterioration inside Tut Ankhamen Tomb (TAT)  
 AUTHOR(S): Louboudy, S. S. ←  
 CORPORATE SOURCE: Bot. & Microbiol., Dept., Fac. of Sci.,  
 Al-Azhar Univ., Cairo, Egypt.  
 SOURCE: Egyptian Journal of Biotechnology, (1998)  
 Vol. 4, pp. 110-123.  
 CODEN: EJBIF7. ISSN: 1110-6093.  
 COUNTRY: EGYPT  
 DOCUMENT TYPE: Journal  
 FILE SEGMENT: CAPLUS  
 OTHER SOURCE: CAPLUS 1999:649978  
 LANGUAGE: English  
 ENTRY DATE: Entered STN: 16 Nov 2001  
 Last Updated on STN: 9 May 2002

AB The purification and properties of L-asparaginase (I)  
 produced by A. niger S-48 TAT, the causal factor of biodeterioration  
 inside the Pharaoh Tutankhamen tomb (TAT), is reported. The  
 purification procedure involved cell-free filtrate preparation (specific  
 activity of 8.92 U/mg protein/mL), fractional precipitation with (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>,  
 (specific activity of 21.05 U/mg protein/mL corresponding to a 2.35-fold  
 purification), dialysis against distilled water followed by dialysis against sucrose  
 crystals, (specific activity of 36.92 U/mg protein/mL, corresponding to a  
 5.7-fold purification) and finally applying a column of Sephadex G-100 (specific  
 activity of 61.0 U/mg protein/mL corresponding to a 6.83-fold  
 purification). The regulatory role of different buffers applied at different pH values  
 revealed that purified I exhibited a maximum specific activity of 62.8  
 U/mg protein/mL in the presence of citrate-phosphate buffer pH 6.6, followed  
 by citrate buffer pH 6.0 (specific activity of 55.46 U/mg  
 protein/mL) and then Tris-HCl buffer pH 7.4 which revealed an obvious decrease  
 in the specific activity (34.16 U/mg protein/mL). By testing purified I

in the presence  
of different substrates, it was found that the highest  
activity was  
obtained by using the most preferable one, i.e., L-  
asparagine, followed by  
L-aspartic acid, L-glutamine, and L-glutamic acid,  
whereas L-arginine,  
L-ornithine, L-threonine and L-citrulline showed  
negligible or inhibitory  
effects toward the purified enzyme activity. Moreover,  
the application of  
different heavy metal cations (in the form of chloride  
salts in addition to  
KCN) as activators and/or inhibitors indicated that  
CaCl<sub>2</sub>, NH<sub>4</sub>Cl, BaCl<sub>2</sub>,  
and MnCl<sub>2</sub> promoted I activity, whereas AlCl<sub>3</sub>, KCN, NiCl<sub>2</sub>,  
ZnCl<sub>2</sub>, FeCl<sub>2</sub>,  
and MgCl<sub>2</sub> had deleterious effects on enzyme activity.  
Purified I was  
tested at different incubation temps., and showed obvious  
activity within  
the temperature range of 22.5-45° with a maximum at 30°.